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Estimates of the Electromagnetic Radiation From Detonation of Conventional Explosives

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Abstract

An order of magnitude model is presented to estimate radiation from detonation of conventional explosives in an attempt to predict frequency bands and signal levels detected by other investigators. An earlier model describing the radiation generated by explosions has been refined to include the contribution of the heat capacity of the detonation products and the temperature dependence of the concentration of ionized particles. Relationships are established between explosions of uncased Composition B, the radiation frequency bands, and the E - and B -field amplitudes as a function of detection distance. The model considers the radiation from particles ionized by passage of the shock wave.

A comparison of the calculated radiation with thermal and background noise estimates shows that the radiation is not detectable above the background radiation even for large explosives at close distances. The fact that radiation has been observed indicates either that the assumptions over-simplify the phenomenon or that the primary mechanism of radiation production has been overlooked. Ionized particles exist 10^6 to 10^8 times longer than the time to accelerate across the shock wave, during which the particles could produce radiation by some other means. Therefore, it is likely that the model greatly underestimates the quantity of radiation produced.

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1. Introduction

1.1 Background

This work is the result of a U.S. Army Research Laboratory (ARL) technology base program to investigate the causes of electromagnetic radiation from the detonation of conventional explosives in support of future combat systems. The overall program objective was to provide the technology to develop a passive, portable, self-contained unit that can use input from multiple sensors to detect, locate, and identify sources of weapons fire and explosives detonation. An estimate of the radio frequency (RF) radiation from the detonation of conventional explosives is presented here, which updates an earlier ARL model [1].

Previous reports [1–3] have addressed several aspects of the problem. Reference [1] provided an order-of-magnitude model based on the thermal effects of explosive materials in producing ionized particles that generate the radiation. Emphasis was on blast effects that had propagated far from the actual detonation. Reference [2] addressed the problem of RF effects on a projectile fuze as it passed directly through a shaped charge explosion. Emphasis was on effects close to the detonation region, and information was provided about the heat capacity of the detonation products and effects of temperature in the explosive region on the concentration of ionized particles. Reference [3] was a literature survey of previously detected RF energy associated with battlefield munitions [4–12].

This report describes refinements of the model of Reference [1] that include consideration of the contribution of the heat capacity of the detonation products and the temperature effect on the concentration of ionized particles. We estimate the characteristics of the RF emission from a detonation of a bare explosive charge about the size of a mortar projectile.

1.2 Summary of Literature Survey

The most prominent results of earlier work are summarized in table 1. Previous investigators detonated explosives including lead azide, RDX (rapidly detonating explosive), Composition B, Tetryl, and trinitrotolul hexogen. Charge sizes from fractions of a gram to 500 tons (5×10^8 grams) were reported. Some radiation was from large caliber Navy projectiles, and other radiation was from bare explosives or metal-encased explosives. Observed radiation frequencies were reported from 1 Hz to 1 GHz. Delay times from initiation of the detonation to observation of the radiation ranged from 80 to 600 μ s.

Two investigators reported *E*-field values at distances close to the detonation. Takakura [5] reports 4×10^{-4} V/m in the frequency band from 6 to 90 MHz at distances of 1m from 0.1 to 0.4 g of lead azide. Using the inverse distance dependence for the radiated *E*-field, we have extrapolated these to

4×10^{-8} V/m at 10 km. Curtis [7] reported 5×10^{-4} V/m from 10 g of RDX at a distance of 6.1 m in the frequency range up to 350 Hz. This extrapolates to 3×10^{-7} V/m at 10 km.

In most cases, the researchers suggested possible physical mechanisms (see table 1) but presented no analytical model. Modeling is difficult because there may be multiple causes. Furthermore, radiation from tube-launched weapons such as small arms, mortar, and artillery may exhibit radiation that is comparable to bomb or projectile detonations because the tube walls conduct charged particles and concentrate the expanding propellant gases in one direction.

Table 1. Frequency bands observed by investigators reviewed.

Investigator	Type of explosive used	Amount of explosive used	Delay / duration of observed signals	Frequency range	Possible cause suggested by authors
Experimental values of frequency ranges					
Trinks	Tube-launched artillery projectiles	None given	—	1–100 kHz	Muzzle flash, ionization of gases near muzzle.
				2 MHz–1GHz	Pulses upon impact at target.
				10 MHz–2GHz	Radiation at detonation from “microsparks” caused by charge equalization at detonation.
Takakura	Lead azide	0.1–0.4 g	80–100 μ s delay	6–90 MHz	Acceleration of electrons ejected by ionization and dipole formation at shock front.
Stuart	Large caliber guns		—	250 MHz–1 GHz	None given, experimental results only.
Curtis	RDX	10 G	2-s delay / 19 s duration	0.5–350 Hz	None given, experimental results only.
Gorshunov et al	50/50 trinitrotolul hexogen	1000–5000 g	—	3 Hz–20 MHz	Electrical charges generated asymmetrically from scattered electrified detonation products
Cook	Composition B	70–1100 g	—	Below 10 kHz	Gaseous detonation products form a plasma at surface of gas cloud from ionization by passing through earth’s electric field. Gas cloud discharges on contact with ground.

Table 1 (cont'd). Frequency bands observed by investigators reviewed.

Investigator	Type of explosive used	Amount of explosive used	Delay / duration of observed signals	Frequency range	Possible cause suggested by authors
Experimental values of frequency ranges					
Wouters	None given	1,300 g 500 ton (= 4.5×10^8 g)	None explicitly given 8-ms duration (1.3 kg) 32-ms duration (500 ton)	—	Blast temperature ionizes detonation products and ambient air and produces a plasma.
van Lint	Bare spheres to metal-cased bombs	10–345,000 g (bare spheres to metal-encased bombs)	100–200- μ s delay	50 MHz–1 GHz	Separation of charge at interface of explosion products and air to form a vertical dipole moment, with asymmetry induced by reflection of shock wave from ground. Electric sparks from explosion products interacting with casing fragments.
Andersen and Long	Bare, plaster-encased, and seeded explosives Tetryl, Composition B	20–1,087 g	300–600- μ s delay	Less than 600 kHz	Detonation ionizes detonation products, which transfer charge by friction to inert casing particles and fragments.

2. Theoretical Model

2.1 The Detonation

We present an order-of-magnitude estimate of the electromagnetic radiation from a detonation or explosion. We use the term “explosion” in the sense described by Wilfred Baker [13] as “a process by which a pressure wave of finite amplitude is generated in air by a rapid release of energy. Some widely different types of energy sources can produce such pressure waves and thus be classified as ‘explosives’ according to our definition....”

“Regardless of the source of the initial finite pressure disturbance, the properties of air as a compressible gas will cause the front of this disturbance to steepen as it passes through the air...until it exhibits nearly discontinuous increases in pressure, density, and temperature. The resulting shock front moves supersonically, faster than sound speed in the air ahead of it. The air particles are also accelerated by the passage of the shock front, producing a net particle velocity in the direction of travel of the front.” This report uses the terms “detonation” and “explosion” interchangeably.

The model is used to estimate the frequency bandwidth of the radiation produced and the electric (E) and magnetic (B) field amplitudes versus distance from the explosion. Results are presented at 10 km from the explosion. The reader can obtain results at other distances by using the formulas that are provided. The model’s results are a function of the temperature within an explosive region. We establish this temperature dependence by assuming an idealized detonation model, is shown in figure 1. The assumptions used in the estimate are italicized.

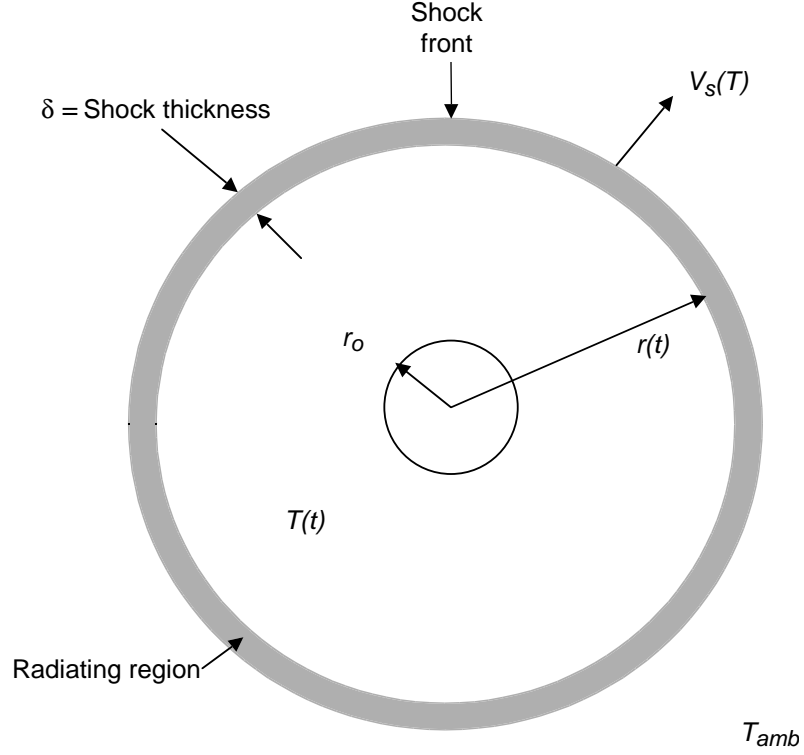
Some of these assumptions are

- *The undetonated projectile occupies a spherical volume defined by radius r_0 .*
- *The detonation is initiated at $t=0$ at the center of a spherical explosive charge.*
- *After initiation of the explosion, a detonation wave propagates outward through the explosive charge with a velocity of $10 \text{ mm}/\mu\text{s} = 10^4 \text{ m/s}$ [14].*
- *At completion of the detonation at $t=t_0$ ($3 \mu\text{s}$ in this example), the detonation wave has expanded to radius r_0 , and all the undetonated explosive has been converted to detonation products with the release of thermal energy into the volume of radius r_0 [1]. We neglect the portion of energy that is converted into kinetic energy of bomb fragments, explosive products, and optical and acoustic radiation.*
- *No expansion occurs for $t < t_0$.*

The temperature of the region with $r < r_0$ at $t = t_0$ is given by

$$T(t_0) = \frac{Q}{C_{V/Det \text{ Pr od}}} + T_{amb} \quad (1)$$

Figure 1. Idealized detonation model. (The undetonated charge occupies the spherical region of radius r_0 . After completion of the detonation, a shock wave moves out with a velocity V_s . The thickness of the temperature discontinuity across the shock wave can be calculated by the formulas provided in the text.)



in which $T_{amb} = 300K$ (ambient temperature at sea level), (2)

and $C_{V/det prod}$ is the average heat capacity of the material inside the sphere of radius r_0 .

For $t > t_0$, the shock wave spreads and the temperature of the enlarging spherical region bound by $r(t)$ is determined by the heat capacity of the additional air as well as the detonation products: The energy, temperature, and pressure are distributed uniformly inside the spherical region behind the shock wave.

We assumed that the specific heat capacity of air is constant and equal to the mean value between specific heat capacity at the elevated density and pressure that exists behind the shock wave at t_0 and specific heat capacity at ambient pressure and temperature ahead of the shock wave. This assumption greatly simplifies our estimate of the temperature rise inside the spherical shock region. The errors as large as a factor of 5 introduced by this assumption are negligible compared to errors of several orders of magnitude introduced by the temperature variation of the fraction of particles ionized, as we discuss later.

For $t > t_0$ and $r < r(t)$

$$T(t) = \frac{Q}{C_{V/air}(t) + C_{V/det prod}} + T_{amb} , \quad (3),$$

$$\text{in which } C_{V/air}(t) = \left(\frac{4}{3} \pi (r^3(t) - r_0^3) \right) \rho_{amb} c_{Vs} . \quad (4)$$

Here, ρ_{amb} is the density of air at ambient temperature and pressure (1.23 kg/m^3) and c_{Vs} is the constant-volume specific heat capacity of air at ambient temperature and pressure (717 J/kg-K).

The velocity of the shock wave as a function of temperature is calculated by Zel'dovich and Raizer [15]. The velocity of the shock wave can be described by their results, which we have fitted to the expression (see fig. 2):

$$V_s(t) = 5.5T^{0.76} \text{ m/s for } (293 \text{ K} < T < 15,000 \text{ K}) . \quad (5)$$

Once we know the initial velocity of the shock wave, $V_s(t_0)$, we can obtain the radius at the subsequent times by using the following equations:

$$r(t_i) = r(t_{i-1}) + V_s(t_{i-1}) \times (t_i - t_{i-1}) \quad (6)$$

in which

$$t_i = t_{i-1} + \Delta t \text{ for } i \geq 1$$

with $\Delta t = 10^{-6} \text{ s}$.

We must determine the value of $C_{v/det \text{ prod}}$ in equation (3) as a function of temperature and pressure.

2.2 Temperature and Pressure Dependence of Heat Capacity of Detonation Products

Composition B explosive consists of 63 percent RDX, 36 percent Trinitrotoluene (dynamite) (TNT), and 1 percent paraffin filler. *The model assumes that the detonation takes place in a vacuum, i.e., in the absence of air. Thus, the program does not include further combustion of the detonation products in air.* The combustion product gases expand outward adiabatically after the completion of detonation. The detonation products of Composition B were obtained from the Cheetah computer program [16] that used the applicable chemistry. The detonation products obtained from the Cheetah model in concentrations greater than 1 mole of detonation product per kilogram of explosive are by weight, 24.1 percent H_2O , 36.4 percent N_2 , 21.9 percent CO_2 , 5.2 percent CO , and 12.1 percent solid carbon.

The specific heat capacity of the detonation product gases depends on temperature and pressure [17], [18]. Temperature dependence [17] is shown in figure 3. The total heat capacity, calculated from figure 3 and the relative concentrations, varies from 0.26 cal/gm-K at 300 K to 0.39 cal/gm-K at 3500 K. The pressure dependence is given in [17]. *The specific heat capacity of the*

Figure 2. Shock wave velocity as a function of temperature behind the shock wave.

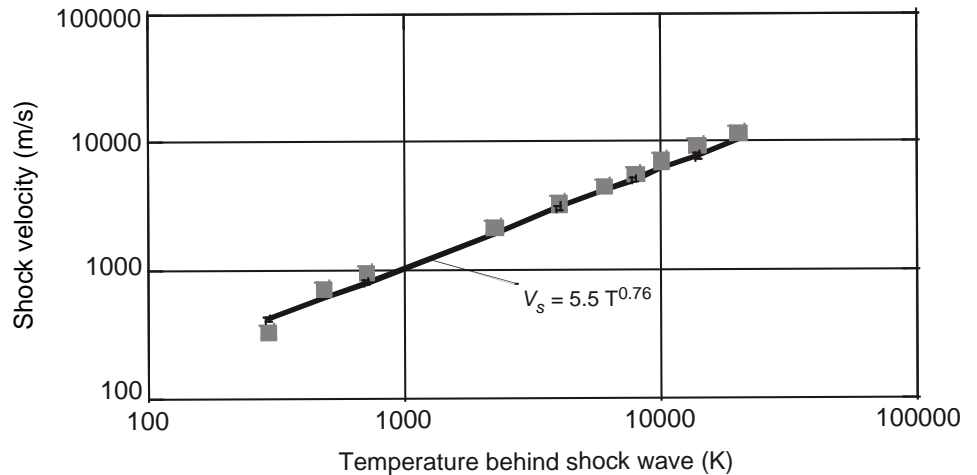
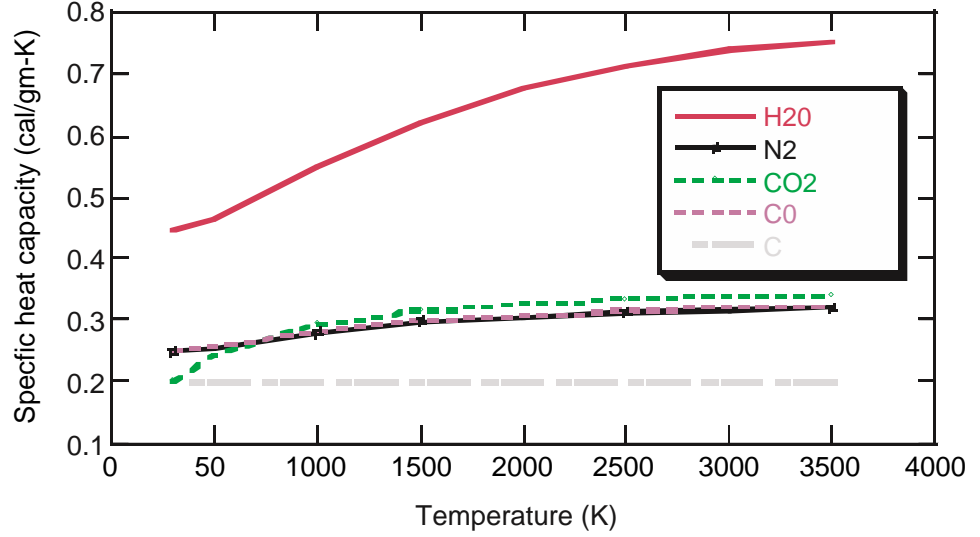


Figure 3. Temperature dependence of the specific heat capacity of the detonation products of Composition B. (The specific heat capacity of each of the five highest concentration detonation products of Composition B is shown.)



detonation products varies adiabatically from 0.210 cal/gm-K at 1 atm to 0.587 cal/gm-k at 24×10^4 atm. In the calculation, we use the mean value of 0.40 over these temperature and pressure ranges.

The pressure and temperature variations on the specific heat capacity of air vary from 0.1719 cal/gm-K at 1 atm and 300 K to 0.315 cal/gm-K at 100 atm and 3000 K. *We use the mean value over this temperature and pressure range, 0.243 cal/gm-K.* This could result in values that are too large or too small by 30 percent over the pressure and temperature extremes.

These assumed values for the specific heat capacities of the detonation products and ambient air lead to errors that are negligible compared to the errors arising from the temperature dependence of the concentration of ionized particles.

2.3 Ionization Fraction

A previous report [2] showed that only a very small fraction of the particles are ionized. We therefore need a method that accounts for the ionization fraction of the radiation. The Hilsenrath and Klein calculation [19] of the ionization fraction of air as a function of temperature is shown in figure 4. One sees that the ionization fraction is very sensitive to changes in temperature below 15,000 K. We have interpolated the lower temperature ionization fraction results from Hilsenrath and Klein in figure 4. The interpolation formula is

$$\text{Log}_{10} \text{IonizationFraction} = \text{Log}_{10} f_i = (-30.566) + (0.011267) * T - (1.3664 \times 10^{-6}) * T^2 \quad (7)$$

Table 2 shows the ionization of various ion species in air at 3500 K and 2500 atmospheres [19] at the high temperatures and pressures that could occur during an explosion. Since air is composed primarily of N_2 and O_2 , and since the dominant detonation products of Comp B explosive are H_2O , N_2 , C, O_2 , and CO_2 , one might expect ionization fraction values of the same order of magnitude as air, which are calculated from equation 7 and plotted in figure 4. We see, however, that the tabulated results from table 2 are quite

Figure 4. Curve fit to ionization fraction, IF, of air as a function of temperature. (The logarithm of the ionization fraction data from Reference 19 is plotted versus temperature of the air. The line obtained from the curve fit equation is also shown.)

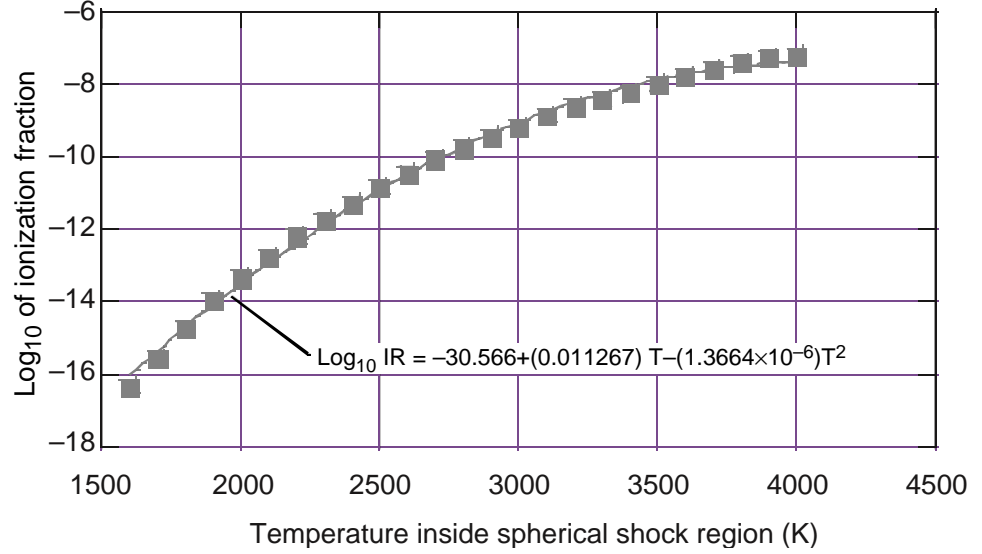


Table 2. Ionization of air at 3500 K and 2500 atm.

Species	Ionization fraction	Trend with increasing pressure
NO ⁺	3.1×10^{-8}	decreasing
O ₂ ⁺	1.5×10^{-11}	decreasing
N ₂ ⁺	2.3×10^{-15}	decreasing
O ⁺	2.5×10^{-15}	decreasing
CO ⁺	1.1×10^{-17}	decreasing
E ⁺	1.0×10^{-8}	decreasing
O ⁺	2.0×10^{-8}	increasing
C ₂ ⁺	1.1×10^{15}	increasing

different from the values for air at 3500 K in figure 4. *Nevertheless, we assume that we can use equation 7 for determining the ionization fraction of the detonation products. We further assume that (a) the detonation products and the air are at most singly ionized, and that (b) the electron concentration, e^- , is equal to the ion concentration.*

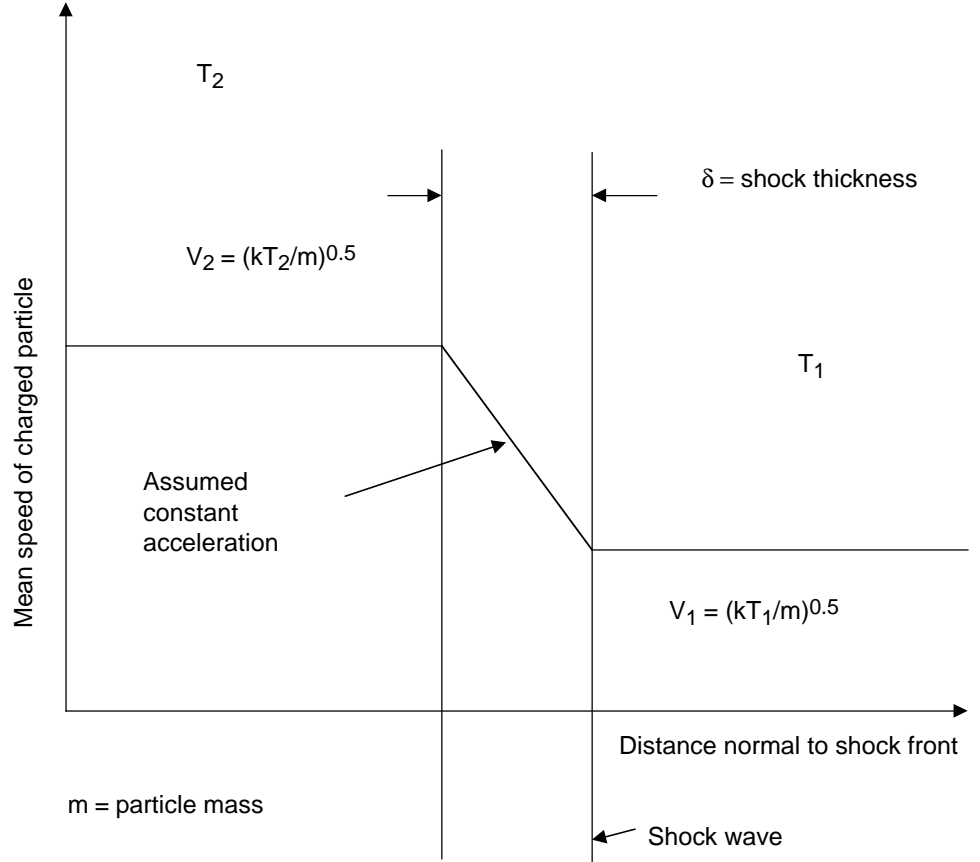
2.4 Radiation From Particle Acceleration Across a Shock Wave

Radiation is caused by the acceleration of charged particles because of the passage of the shock wave. Additional assumptions are as follow: *All the acceleration takes place over the thickness of the shock wave, which we shall show is on the order of 10^{-8} m. We apply the equipartition theorem [20] to provide a relationship for the mean particle speed V . Ideal gas behavior applies, and the air before and after the shock wave is in thermodynamic equilibrium.*

Since we are considering only the radial component of the electron's velocity, the problem is one-dimensional, with the result that the speed change, ΔV , across the shock (neglecting direction changes) and the average speed across the shock, \bar{V} , as shown in figure 5, are given respectively by

$$\Delta V \approx V(t) - V(T_{amb}) = \sqrt{\frac{k}{m}} (\sqrt{T} - \sqrt{T_{amb}}) , \quad (8)$$

Figure 5. Change in mean speed of charged particles caused by temperature change across the shock wave for the one-dimensional case considered in the analysis.



in which k is the Boltzmann constant, 1.38×10^{-23} J/K.

$$\bar{V} \approx \frac{V(T) + V(T_{amb})}{2} = \sqrt{\frac{k}{m}} \left(\frac{\sqrt{T} + \sqrt{T_{amb}}}{2} \right), \quad (9)$$

in which $V(T)$ is the average radial velocity for $r > r_0$.

The shock wave Mach number is defined by

$$M(t) \equiv \frac{V_s(T(t))}{V_{amb}} \quad (10)$$

in which V_{amb} , the speed of sound at ambient conditions, = 340 m/s.

The mean free path λ , which is the average distance that particles travel between collisions [21], is

$$\lambda \approx \frac{16}{5} \frac{\nu}{\sqrt{2\pi R T_{amb}}} \quad (11)$$

in which ν is the kinematic viscosity of air at ambient conditions (14×10^{-6} m²/s) and R is the ideal gas constant of air (287 J/kg-K).

The shock thickness $\delta(t)$ is given by Kogan [21]

$$\delta(t) \approx \frac{\gamma - 1}{\gamma + 1} M(t) \lambda \approx 10^{-8} M(t) \quad (12)$$

in which γ is the ratio of the constant-pressure and constant-volume specific heat capacities of air, which we have assumed to be constant over the range of temperatures and pressures behind the shock.

The volume of air in the thin shock wave is

$$\tau_w = 4\pi r^2(t) \delta(t) . \quad (13)$$

With the assumption of constant-volume and constant-density conditions, the pressure rise produced is the bomb-yield energy divided by the volume of the spherical shock region, so that the pressure is

$$p(t) \approx p_{amb} + \frac{Q}{\frac{4}{3}\pi r^3(t)} \quad (14)$$

in which p_{amb} is ambient pressure, $1.01 \times 10^5 \text{ N/m}^2$.

During these conditions, the number of moles of air in the region in the shock wave is given by [22]

$$n(t) = \frac{p(t) \tau_w(t)}{RT(t)} . \quad (15)$$

Now consider a charged particle that accelerates with an acceleration $a(t)$ across the shock front for a short time Δt and continues moving in the same direction with the new velocity. The time of transit of the charged particle across the shock wave is

$$\Delta t \approx \frac{\delta}{\bar{V}} \quad (16)$$

and the acceleration of the charged particle is given by

$$a \approx \frac{\Delta V}{\Delta t} = \bar{V} \frac{\Delta V}{\delta} . \quad (17)$$

We assume that the radiation is more or less uniformly distributed over a band starting at 0 Hz and extending to $1/\Delta t$ [23], given by

$$\Delta f \approx \frac{1}{\Delta t} . \quad (18),$$

and that from equation (17), we see that

$$\Delta f(t) \approx \frac{\bar{V}}{\delta(t)} . \quad (19)$$

The time-average total power per singly ionized particle, P , integrated over all directions, is given by [24]

$$P = \frac{e^2 a^2}{6\pi\epsilon_0 c^3} = \frac{\mu_0 e^2 a^2}{c 6\pi} , \quad (20)$$

in which e is the electron charge ($1.6 \times 10^{-19} \text{ C}$); μ_0 , the permeability of free space (taken as equal to the permeability of air), is $1.26 \text{ } \mu\text{H/m}$; and ϵ_0 , the permittivity of free space (taken as equal to the permittivity of air), is 8.85 pF/m .

The total power radiated for n moles of singly ionized particles is

$$P_{total} \approx f_i n A P . \quad (21)$$

in which A is the Avogadro's number, 6.02×10^{23} particles/mole, and f_i is the fraction of particles ionized from equation (7).

The average E -field per cycle produced in the far field per particle is given by [25]

$$E = \frac{1}{r} \sqrt{\frac{P \mu_0 c}{2\pi}} \quad (22)$$

Using equations 20, 21, and 22, we obtain the following formulas for the total E -field and B -field:

$$E_{total} \approx \left(\frac{1}{r} \right) \left(\sqrt{\frac{f_i (nA) \left(\frac{e^2 2a^2}{4\pi\epsilon_0^3} \right) \mu_0 c}{2\pi}} \right) \approx \frac{\mu_0 \sqrt{3}}{2\pi r} (ea) \sqrt{f_i n a} \quad (23)$$

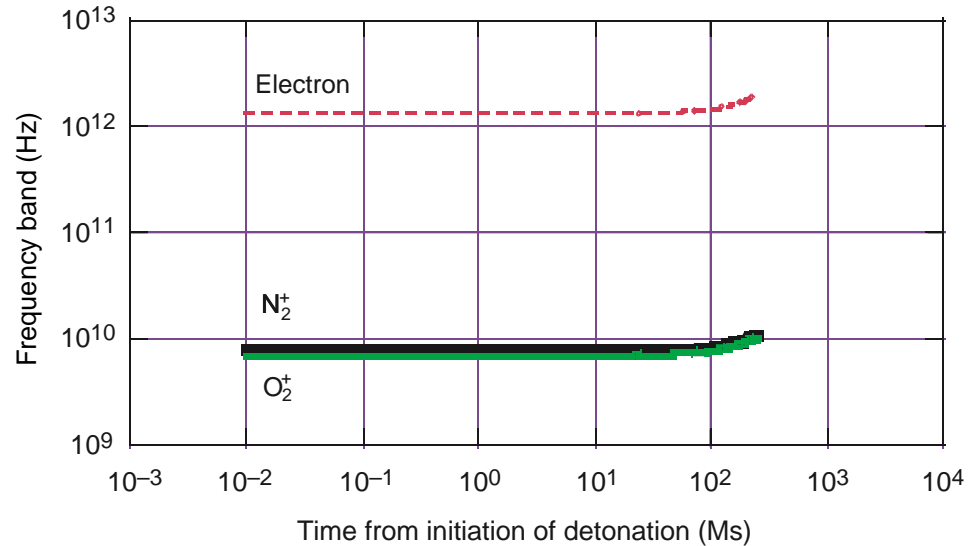
$$B_{total} \approx \frac{E_{total}}{c} (\text{Wb/m}^2) . \quad (24)$$

Calculations have been performed for the case of an explosion having a yield that approximates the energy from a mortar projectile that is 25 MJ.

The calculated frequency bands shown in figure 6 include the observed frequency ranges of 6 to 250 MHz to 2 GHz, as shown in table 1. The electron frequency band is 1.4 THz, which is too high to be observed with RF, non-line-of-sight equipment. Figure 6 shows that the upper limits of the frequency bandwidths for N_2 and O_2 ions are about 8 to 9 GHz.

Figure 7 is the calculated E -field amplitude at a range of 10 km versus time from initiation of detonation for the O_2 ions, N_2 ions, and electrons. The E -field amplitude depends on the mass of the particles through the acceleration (eqs 17 and 23), the velocity change and average velocities (eqs 8 and

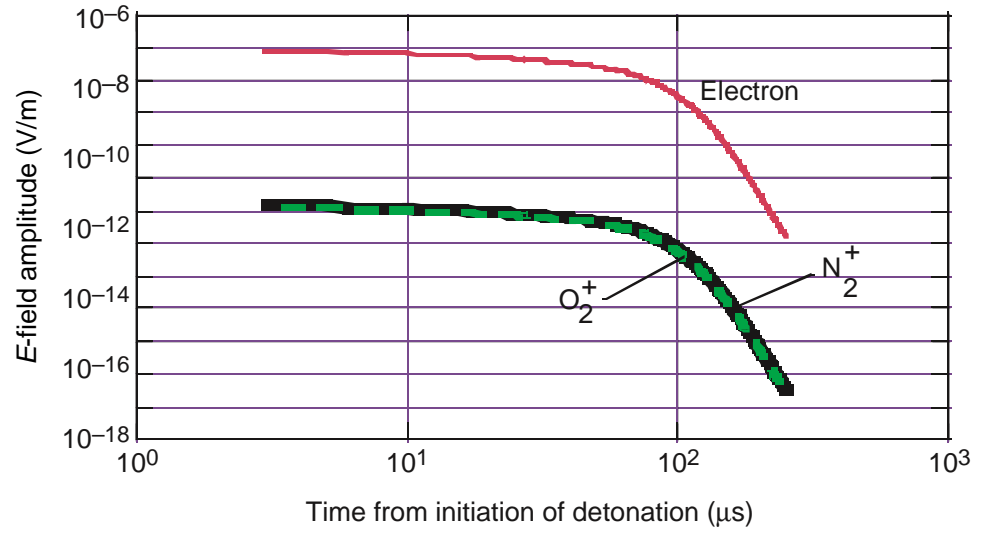
Figure 6. Calculated frequency band of radiation from electron, O_2^+ , and N_2^+ ions formed when the shock wave from the detonation passes through ambient air. (The frequency band in volts/meter is plotted versus time in microseconds from initiation of detonation at 10 km.)



9). If equations (8) and (9) are substituted into equation (17), and the resulting acceleration is substituted into equation (23), the resulting expression shows that the E -field amplitude is inversely proportional to the masses of the particles. Thus, the radiation from the electrons should be more detectable than the radiation from the heavier O_2^+ and N_2^+ .

If we arbitrarily assume that the detection threshold is 10^{-7} V/m, then from figure 7, the electron's radiation is detectable at 10 km for 100 μ s, but the radiation from the O_2^+ and N_2^+ ions is not detectable. At 10 m from the explosion, the signal from the ions would increase to 2×10^{-7} V/m, and would remain above the detection threshold for about 3 μ s.

Figure 7. Calculated electric field amplitude of radiation from electron, O_2^+ , and N_2^+ ions formed when the shock wave from the detonation passes through ambient air. (The E -field amplitude in volts/meter is plotted versus time in microseconds from initiation of the detonation at 10 km.)



3. Consequences of the Model

3.1 Effect of Explosive Mass

The model shows that the mass of the undetonated explosive affects both the amplitude of the radiation and the delay time from initiation of the explosion to completion of the detonation.

Equation (23) shows that the E -field amplitude is proportional to the square root of the number of moles of ionized particles, which *we assume is proportional to the mass of the undetonated explosive*. Thus, the model leads us to expect that the amplitude ratio should be proportional to the square root of the explosive mass.

The delay time from initiation to completion of the detonation is approximately the time for the detonation wave to propagate at constant velocity, V_{det} from the center of the spherical explosive (point of initiation) to the surface of the explosive. From reference [3], the ratio of the delay times for two spherical explosives of uniform density but different masses should be proportional to the ratio of the cube roots of their masses.

3.2 Radiation From Recombination of Ions with Electrons

Heretofore, we have discussed the radiation caused by particles ionized during their passage through the shock region; we have not considered that the ions eventually re-combine with electrons, producing further radiation. Table 3 [26] gives the ionization energies and frequencies of the detonation products N_2 and CO_2 .

The frequencies are in the terahertz region, with wavelengths in the ultra-violet spectrum, and therefore are not useful for transmission in non-line-of-sight scenarios.

3.3 Detectability of RF Signals From Detonation of Conventional Explosives

The detectability of the radiation depends on the thermal noise, background radiation noise, and the noise inherent in the sensor and data acquisition system. The examination of particular instrumentation systems and background radiation noise is beyond the scope of this report; however, we can make some general observations about the thermal noise. The thermal noise power is given by [27]

$$S_{noise} = kT_{amb}\Delta f \quad (25)$$

Table 3. Radiation from recombination of detonation products.

Detonation product	Ionization energy (e-V)	Ionization energy (J)	Frequency (Hz)	Wavelength (Å)
N_2	15.5	2.48×10^{-18}	3.73×10^{15}	800
CO_2	14.4	2.30×10^{-18}	4.0×10^{15}	750

in which T_{amb} is the absolute temperature of the circuits in K, here taken to be 300 K, and Δf is the frequency bandwidth of the signal in hertz. The calculated values of Δf and E from figures 6 and 7 are given in table 4, with the noise power calculated from equation (25).

The power P_{avg} received by a 1-m² antenna far from the source can be estimated from the plane wave relationship [28], via E -field values obtained from figure 7:

$$P_{avg}(\text{watts}) = \left[\frac{1}{2} c \epsilon_0 E_{peak}^2 \right] A_{eff} (m^2) = \left[0.00133 E_{peak}^2 \right] A_{eff} , \quad (26)$$

c and ϵ_0 were defined previously.

Table 4 shows that the average power received is far less than the $kT\Delta f$ noise as close as 1 m to the detonation. The electron, which is the most detectable particle, is 11 dB below the noise, even at 1 m, and is 91 dB below noise at 10 km. Even if an explosive 10,000 times larger than the mortar (equivalent to a 4000-lb bomb) is used, only the electron radiation is detectable above the noise at close distances.

Using E -field amplitudes and frequency bands from figures 6 and 7, we can calculate the signal level relative to the noise at 1 m and 10 km from the detonation. The data are then extrapolated to an explosive having 10,000 times more mass.

These calculations have shown that the wide-band signals can be expected from radiation caused by ionized particles accelerating across the shock wave; however, the signals are not detectable above $kT\Delta f$ noise for normal sized explosives at useful detection ranges. Thus, the assumptions have oversimplified the phenomenon, or the model has not established the mechanism responsible for the signals measured by previous investigators (see table 1 and sect. 2).

Table 4. Detectability of estimated radiation from shock wave.

Particle	Calculated E -field (V/m)	Δf (Hz)	Noise power (W)	Calculated Average power at receiver (W)	Detectability (dB)
0.191 kg explosive (mortar)					
At 1 m					
Electron	6×10^{-4}	1.6×10^{12}	6×10^{-9}	4.8×10^{-10}	-11
N ₂ Ion	1×10^{-8}	9×10^9	3.7×10^{-11}	1.3×10^{-19}	-84
O ₂ Ion	9×10^{-9}	8×10^9	3.3×10^{-11}	1.07×10^{-19}	-85
At 10 km					
Electron	6×10^{-8}	1.6×10^{12}	6×10^{-9}	4.8×10^{-18}	-91
N ₂ Ion	1×10^{-12}	9×10^9	3.7×10^{-11}	1.3×10^{-27}	-164
O ₂ Ion	9×10^{-13}	8×10^9	3.3×10^{-11}	1.07×10^{-27}	-165

3.4 Discussion

The calculations and order-of-magnitude model address the radiation produced by explosions that occur in bomb and projectile detonations. The bare explosive (not the explosive encased by the bomb or projectile shell) is considered. The mechanism of radiation analyzed was radiation from particles ionized by passage of the shock wave. The most common charged particles from the atmosphere are the O_2^+ ions and N_2^+ ions and the electrons. The calculations that were presented for each type of particle are the radiation bandwidth, time-average E -field and B -field amplitudes.

The values of bandwidth measured by experimenters (see table 1) ranged from 1 MHz to 1 GHz. The values from the model extend from 8 GHz for the ions to terahertz for the electrons.

The heat capacity of the detonation products and of the ambient air determine the temperature of the region of high temperature, pressure, and density behind the shock wave. The concentration of ionized particles strongly depends on the temperature behind the spherical shock wave. It decreases very steeply as the temperature drops during the shock wave expansion. An estimate was made of the dependence of the ion concentration on temperature. Relationships were established between explosive mass of uncased Composition B, the radiation frequency bands, and the E - and B -field amplitudes as a function of detection distance. At a range of 10 km, radiation from free electrons caused by a mortar-sized blast is expected in the frequency band up to 1.6 THz with amplitudes above 0.01 microvolt/m for the first 100 μ s from detonation. Oxygen and nitrogen ions produce radiation in the frequency band up to 9 GHz with amplitudes above 10 picovolt/m for 3 μ s after detonation.

A comparison of the calculated radiation with thermal noise estimates shows that most of the radiation is not detectable above the noise even for large explosives at close distances. The fact that signals have been observed suggests either that the assumptions over-simplify the phenomenon or that the radiation detected must be caused by other mechanisms.

The times for the ions and electrons to cross the shock wave are the reciprocals of the frequency bands from figure 6: 10^{-10} s for the ions and 10^{-12} s for the electrons. The particles exist in the charged state for the duration of the explosive event, which from figures 6 and 7, is nominally 100 μ s or 10^{-4} s. This is 10^6 times longer than the shock transit time for the ions and 10^8 times longer for the electrons. During this much longer time, the charged particles could be producing radiation by some other mechanism. We therefore believe that the model greatly underestimates the quantity of radiation produced.

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